

Page 107

Please amend the paragraph at page 107, lines 3-12 as follows:

Compound 69 72 can be produced in the following manner: compound 69 which is easily in accordance with the known methods (U=hydroxyl group: for example, *J. Org. Chem.* 1992, 57, 5680-5686., U=thiol group: for example, *J. Heterocycle. Chem.* 1990, 27, 567., U=amino group: for example, *Synthesis* 1987, 1124.) is iodized or brominated at 4-position of pyrazole to obtain compound 70; trimethylsilyl acetylene is coupled to this compound 70 by Sonogashira coupling and detrimethylsilylation is conducted to obtain compound 71; and then compound 71 is aromatized and deprotected.

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Please amend the paragraph at page 301, lines 12-21 as follows:

To a solution of 31.3 mg of 3-[(*E*)-2-(4-fluorophenyl)-vinyl]-4-(2-hydroxyethoxy)-1*H*-indazole-5-carboxylic acid ethyl ester obtained by Example 349-*j* in 0.7 mL of tetrahydrofuran were added 0.3 mL of ethanol and 0.2 mL of 5N sodium hydroxide aqueous solution, and stirred at 70°C for 1 hour and 20 minutes. After cooling on ice, the solution was neutralized with 2N hydrochloric acid under ice cooling, and the precipitated crystals were collected by filtration, and dried under reduced pressure, to afford 25 mg of the title compound as yellow crude crystals.

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Please amend the paragraph at page 506, lines 15-26 as follows:

Example 1054

10/509,795

Application No.: Not Yet Assigned

Docket No.: 0425-1154PUS1

1-Ethyl-3-{6-fluoro-3-[(*E*)-2-(4-fluorophenyl)-vinyl]-1*H*-indazol-5-yl}-urea

10 mg of 6-fluoro-3-[(*E*)-2-(4-fluorophenyl)-vinyl]-1-trityl-1*H*-indazol-5-ylamine

obtained by Production example 182 and 3 μ l of ethyl isocyanate was dissolved in 5 mL of chloroform, and heated at reflux for 2 hours. The reaction solution was allowed to cool to room temperature, added with 0.5 mL of trifluoroacetic acid, stirred at room temperature for 30 minutes, and the reaction mixture was purified and separated by LC-MS, to afford 0.67 mg of the title compound as pale yellow powder.

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Please amend the paragraph at page 526, lines 5-15 as follows:

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Example 1146

7-Fluoro-3-[(*E*)-2-(4-fluorophenyl)-vinyl]-5-[5-(piperidin-1-yl)methyl-4*H*-[1,2,4]triazol-3-yl]-1*H*-indazole

13 mg piperidin-1-yl-acetic acid hydrazide produced by Production example 1139 and 15 mg of 7-fluoro-3-[(*E*)-2-(4-fluorophenyl)-vinyl]-1*H*-indazole-5-carboxymidic acid ethyl ester hydrochloride obtained by Example 383 were dissolved in 1 mL of butanol, and added with 30 μ l of triethylamine. After stirring at 105°C for 8 hours, the solution was purification by LC-MS, to afford 2.08 mg of the title compound.

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Please amend the paragraph at page 577, lines 14-18 as follows:

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$^1\text{H-NMR}$ (400MHz, DMSO-D₆) δ 1.17 (2H, dd, J=4.4, 7.6Hz), 1.43 (2H, dd, J=4.8, 7.6Hz), 3.96